DERs needed for MRID Studies for Momfluorothrin (2014 EPA Momofluorothrin: Human Health Risk Assessment for a Proposed Use of the New Active Ingredient In Indoor Residential Settings and Outdoors in Commercial/Residential Settings)

https://www.regulations.gov/document?D=EPA-HQ-OPP-2015-0752-0008

MRID number	Study type
49019993	Acute oral toxicity rat
<mark>49019995</mark>	Acute dermal toxicity rat
<mark>49019997</mark>	Acute inhalation toxicity rat
<mark>49019999</mark>	Acute eye irritation rabbit
<mark>49020002</mark>	Acute dermal irritation rabbit
49020004	Skin sensitization guinea pig
<mark>49020006</mark> *	90-day oral toxicity rat
<mark>49020008</mark> *	90-day oral toxicity dog
<mark>49020010</mark> *	28-day dermal toxicity rat
<mark>49020012</mark> *	28-day inhalation toxicity rat
<mark>49020014</mark> *	Prenatal developmental toxicity rabbit
<mark>49020016</mark> *	Prenatal developmental toxicity rat
<mark>49020018</mark> *	Reproduction and fertility effects rat
<mark>49020020</mark> *	Chronic toxicity rat
<mark>49020022</mark> *	Carcinogenicity mouse
<mark>49020024</mark> *	Carcinogenicity rat
<mark>49020026</mark> *	Reverse mutation
<mark>49020028</mark> *	In vitro mammalian gene mutation
<mark>49020030</mark> *	In vitro mammalian chromosome aberration
<mark>49020032</mark> *	Mammalian micronucleus
<mark>49020034</mark> *	Unscheduled DNA synthesis
49020036*, 49020038	Acute neurotoxicity rat
<mark>49020040</mark> *	Subchronic neutoroxicity screening battery rat
<mark>49020042</mark> *	Metabolism and pharmacokinetics rat
<mark>49020044</mark> *	Metabolism and pharmacokinetics rat
<mark>49020046</mark>	Dermal penetration in vitro
<mark>49020048</mark> *	Immunotoxicity rat

The studies forwarded in the August 29 response are highlighted.

The studies for which DERs are included are marked with an asterisk (*).

DER's for the following studies will be forwarded in a future response:

MRID 49019993, 49019995, 49019997 & 49019999 MRID 49020002 & 49020004, 49020038 & 49020046